Controlled substance	Drug code	Schedule
Meperidine	9230	II.
Meperidine intermediate-B	9233	II.
Methadone	9250	II.
Dextropropoxyphene, bulk (non-dosage forms)	9273	II.
Morphine	9300	II.
Thebaine	9333	II.
Oxymorphone	9652	II.
Alfentanil	9737	II.
Remifentanil	9739	II.
Sufentanil	9740	II.
Carfentanil	9743	II.
Tapentadol	9780	II.
Fentanyl	9801	II.

The company plans to manufacture bulk controlled substances for use in product development of analytical reference standards, for distribution to its customers.

Dated: December 20, 2016.

Louis J. Milione,

Assistant Administrator.

[FR Doc. 2016–31285 Filed 12–27–16; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Importer of Controlled Substances Registration

ACTION: Notice of registration.

SUMMARY: Registrants listed below have applied for and been granted registration by the Drug Enforcement Administration (DEA) as importers of various classes of schedule I or II controlled substances.

SUPPLEMENTARY INFORMATION:

The companies listed below applied to be registered as importers of various basic classes of controlled substances. Information on previously published notices is listed in the table below. No comments or objections were submitted and no requests for hearing were submitted for these notices.

Company	FR docket	Published
Rhodes Technologies	81 FR 46956	July 19, 2016.
Bellwyck Clinical Services	81 FR 54603	August 16, 2016.
Cerilliant Corporation	81 FR 57933	August 24, 2016.
Noramco, Inc	81 FR 57932	August 24, 2016.
Cody Laboratories, Inc	81 FR 54602	August 16, 2016.
AMRI Rensselaer, Inc	81 FR 54603	August 16, 2016.
ALMAC Clinical Services Incorp (ACSI)	81 FR 54602	August 16, 2016.
Fresenius Kabi USA, LLC	81 FR 54601	August 16, 2016.
Akorn, Inc	81 FR 57935	August 24, 2016.
Actavis Laboratories FL, Inc	81 FR 54602	August 16, 2016.
Unither Manufacturing LLC	81 FR 61250	September 6, 2016.
Cambrex Charles City	81 FR 63222	September 14, 2016.
United States Pharmacopeial Convention	81 FR 63220	September 14, 2016.
R & D Systems, Inc	81 FR 64509	September 20, 2016.
Catalent CTS, LLC	81 FR 66081	September 26, 2016.

The DEA has considered the factors in 21 U.S.C. 823, 952(a) and 958(a) and determined that the registration of the listed registrants to import the applicable basic classes of schedule I or II controlled substances is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971. The DEA investigated each company's maintenance of effective controls against diversion by inspecting and testing each company's physical security systems, verifying each company's compliance with state and local laws, and reviewing each company's background and history.

Therefore, pursuant to 21 U.S.C. 952(a) and 958(a), and in accordance with 21 CFR 1301.34, the DEA has granted a registration as an importer for schedule I or II controlled substances to the above listed persons.

Dated: December 19, 2016.

Louis J. Milione,

Assistant Administrator.

[FR Doc. 2016–31273 Filed 12–27–16; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Bulk Manufacturer of Controlled Substances Application: Johnson Matthey Inc.

ACTION: Notice of application.

DATES: Registered bulk manufacturers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration in accordance with 21 CFR 1301.33(a) on or before February 27, 2017.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DRW, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: The Attorney General has delegated her authority under the Controlled Substances Act to the Administrator of the Drug Enforcement Administration (DEA), 28 CFR 0.100(b). Authority to exercise all necessary functions with

respect to the promulgation and implementation of 21 CFR part 1301, incident to the registration of manufacturers, distributors, dispensers, importers, and exporters of controlled substances (other than final orders in connection with suspension, denial, or revocation of registration) has been redelegated to the Assistant Administrator of the DEA Diversion Control Division (''Assistant Administrator'') pursuant to section 7 of 28 CFR part 0, appendix to subpart R.

In accordance with 21 CFR 1301.33(a), this is notice that on September 5, 2016, Johnson Matthey Inc., Pharmaceuticals Materials, 900 River Road, Conshohocken, Pennsylvania 19428, applied to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
Gamma Hydroxybutyric Acid	2010	I
Amphetamine	1100	II
Methylphenidate	1724	II
Codeine	9050	II
Oxycodone	9143	II
Diphenoxylate	9170	II
Hydrocodone	9193	II
Meperidine	9230	II
Methadone	9250	II
Methadone intermediate	9254	II
Morphine	9300	II
Thebaine	9333	II
Opium tincture	9630	II

The company plans to manufacture the listed controlled substances in bulk for distribution and sale to its customers. Thebaine (9333) will be used to manufacture other controlled substances for sale in bulk to its customers.

Dated: December 20, 2016.

Louis J. Milione,

Assistant Administrator.

[FR Doc. 2016–31283 Filed 12–27–16; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF LABOR

Office of the Secretary

Agency Information Collection Activities; Submission for OMB Review; Comment Request; Request for State or Federal Workers' Compensation Information

ACTION: Notice.

SUMMARY: The Department of Labor (DOL) is submitting the Office of Workers' Compensation Programs (OWCP) sponsored information collection request (ICR) revision titled, ''Request for State or Federal Workers' Compensation Information,'' to the Office of Management and Budget (OMB) for review and approval for use in accordance with the Paperwork Reduction Act (PRA) of 1995 (44 U.S.C. 3501 *et seq.*).

DATES: Submit comments on or before January 27, 2017.

ADDRESSES: A copy of this ICR with applicable supporting documentation; including a description of the likely respondents, proposed frequency of response, and estimated total burden may be obtained free of charge from the *RegInfo.gov* Web site at *http://www.reginfo.gov/public/do/PRAViewICR?ref_nbr=201607-1240-002* (this link will only become active on the day following publication of this notice) or by contacting Michel Smyth by telephone at 202–693–4129 (this is not a toll-free number) or sending an email to *DOL_PRA_PUBLIC@dol.gov.*

Submit comments about this request to the Office of Information and Regulatory Affairs, Attn: OMB Desk Officer for DOL–OWCP, Office of Management and Budget, Room 10235, 725 17th Street NW., Washington, DC 20503, Fax: 202–395–6881 (this is not a toll-free number), email: *OIRA_submission@omb.eop.gov.* Commenters are encouraged, but not required, to send a courtesy copy of any comments to the U.S. Department of Labor-OASAM, Office of the Chief Information Officer, Attn: Compliance Management Program, Room N1301, 200 Constitution Avenue NW., Washington, DC 20210, email: *DOL_PRA_PUBLIC@dol.gov.*

FOR FURTHER INFORMATION CONTACT: Contact Michel Smyth by telephone at 202–693–4129 (this is not a toll-free number) or by email at *DOL_PRA_PUBLIC@dol.gov.*

Authority: 44 U.S.C. 3507(a)(1)(D).

SUPPLEMENTARY INFORMATION: This ICR seeks approval under the PRA for revisions to the Request for State or Federal Workers' Compensation Information (Form CM–905) information collection. Form CM–905 collects information to process a claim under the Black Lung Benefits Act (30 U.S.C. 901 *et seq.*). The information collected helps determine compensation benefits awarded for pneumoconiosis. The information collection has been classified as a revision, because the OWCP proposes to make a series of cosmetic and minor changes to Form CM–905. The changes provide clearer language, so that Federal/State workers' compensation officials clearly understand which portion of the form they should complete and what information to provide. Other changes update the form to reflect current organizational structure within the DOL. For additional substantive information about this ICR, see the related notice published in the **Federal Register** on July 27, 2016 (81 FR 49270).

This information collection is subject to the PRA. A Federal agency generally cannot conduct or sponsor a collection of information, and the public is generally not required to respond to an information collection, unless it is approved by the OMB under the PRA and displays a currently valid OMB Control Number. In addition, notwithstanding any other provisions of law, no person shall generally be subject to penalty for failing to comply with a collection of information that does not display a valid Control Number. *See* 5 CFR 1320.5(a) and 1320.6. The DOL obtains OMB approval for this information collection under Control Number 1240–0032. The current approval for this collection is scheduled to expire on December 31, 2016; however, the DOL notes that existing information collection requirements submitted to the OMB receive a month-to-month extension while they undergo